

# **Product Data Sheet**

### **Anti-NT5C1A Antibody**

Catalog #	Source	Reactivity	Applications		
CPA4928	Rabbit	H, M, R	WB, IH, IF/IC		
Description	Rabl	Rabbit polyclonal antibody to NT5C1A			
Immunogen	KLH-	-conjugated synthetic p	eptide encompassing a sequence within the center		
	regio	on of human NT5C1A. T	he exact sequence is proprietary.		
Purification	The	antibody was purified b	by immunogen affinity chromatography.		
Specificity	Reco	ognizes endogenous lev	els of NT5C1A protein.		
Clonality	Poly	clonal			
Conjugation					
Form	Liqu	id in 0.42% Potassium p	phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,		
	and	0.01% sodium azide.			
Dilution	WB	(1/500 - 1/1000), IH (1/5	0 - 1/100), IF/IC (1/50 - 1/200)		
Gene Symbol	NT5	C1A			
Alternative Na	ames Cyto	solic 5'-nucleotidase 1/	A; cN1A; Cytosolic 5'-nucleotidase IA; cN-I; cN-IA		
Entrez Gene	8462	18 (Human); 230718 (N	louse)		
SwissProt	Q9B	XI3 (Human); A3KFX0 (I	Mouse)		
Storage/Stabi	lity Ship	ped at 4°C. Upon delive	ry aliquot and store at -20°C for one year. Avoid		
	free	ze/thaw cycles.			

Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC-Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, RNAi- RNA interference Species reactivity key: H- Human, M- Mouse, R- Rat, B- Bovine, C- Chicken, D- Dog, G- Goat, Mk- Monkey, P- Pig, Rb-Rabbit, S- Sheep, Z- Zebrafish

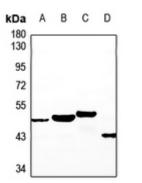
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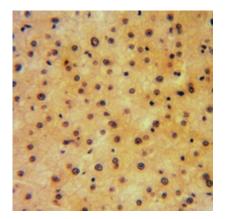




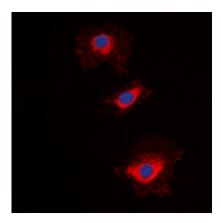
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Western blot analysis of NT5C1A expression in HCT116 (A), COS7 (B), mouse heart (C), PC12 (D) whole cell lysates. (Predicted band size: 41 kD; Observed band size: 41; 45 kD)



Immunohistochemical analysis of NT5C1A staining in human liver cancer formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then incubated with the antibody at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.



Immunofluorescent analysis of NT5C1A staining in HepG2 cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4 °C in a hidified chamber. Cells were washed with PBST and incubated with a DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark. DAPI was used to stain the cell nuclei (blue).

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